



## ΚΥΚΛΟΣ ΣΕΜΙΝΑΡΙΩΝ ΣΤΑΤΙΣΤΙΚΗΣ ΟΚΤΩΒΡΙΟΣ – ΔΕΚΕΜΒΡΙΟΣ 2014

**Brunilda Balliu**

*Department of Medical Statistics  
Leiden University Medical Centre*

### **A Retrospective Likelihood Approach for Efficient Integration of Multiple Omics Factors in Case-Control Association Studies**

ΤΕΤΑΡΤΗ 22/10/2014  
15:00 – 16:00

**ΑΙΘΟΥΣΑ 607, 6<sup>ος</sup> ΟΡΟΦΟΣ,  
ΚΤΙΡΙΟ ΜΕΤΑΠΤΥΧΙΑΚΩΝ ΣΠΟΥΔΩΝ  
(ΕΥΕΛΠΙΔΩΝ & ΛΕΥΚΑΔΟΣ)**

**ΠΕΡΙΛΗΨΗ (ΣΤΑ ΑΓΓΛΙΚΑ)**

Integrative omics, the joint analysis of outcome and multiple types of omics data, such as genomics, epigenomics and transcriptomics data, constitute a promising approach for powerful and biologically relevant association studies. These studies often employ a case-control design, and often include non-omics covariates, such as age and gender, that may modify the underlying omics risk factors. An open question is how to best integrate multiple omics and non-omics information to maximize statistical power in case-control studies that ascertain individuals based on the phenotype. Recent works on integrative omics have used prospective approaches, modeling case-control status conditional on omics and non-omics risk factors. Compared to univariate approaches, jointly analyzing multiple risk factors with a prospective approach increases power in non-ascertained cohorts. However, these prospective approaches often lose power in case-control studies. In this article, we propose a novel statistical method for integrating multiple omics and non-omics factors in case-control association studies. Our method is based on a retrospective likelihood function that models the joint distribution of omics and non-omics factors conditional on case-control status. The new method provides accurate control of Type I error rate and has increased efficiency over prospective approaches in both simulated and real data.



## AUEB STATISTICS SEMINAR SERIES OCTOBER– DECEMBER 2014

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Wednesday 22/10/2014  
**15:00 – 16:00**

**ROOM 607, 6<sup>th</sup> FLOOR,  
POSTGRADUATE STUDIES BUILDING  
(EVELPIDON & LEFKADOS)**

#### **ABSTRACT**

Integrative omics, the joint analysis of outcome and multiple types of omics data, such as genomics, epigenomics and transcriptomics data, constitute a promising approach for powerful and biologically relevant association studies. These studies often employ a case-control design, and often include non-omics covariates, such as age and gender, that may modify the underlying omics risk factors. An open question is how to best integrate multiple omics and non-omics information to maximize statistical power in case-control studies that ascertain individuals based on the phenotype. Recent works on integrative omics have used prospective approaches, modeling case-control status conditional on omics and non-omics risk factors. Compared to univariate approaches, jointly analyzing multiple risk factors with a prospective approach increases power in non-ascertained cohorts. However, these prospective approaches often lose power in case-control studies. In this article, we propose a novel statistical method for integrating multiple omics and non-omics factors in case-control association studies. Our method is based on a retrospective likelihood function that models the joint distribution of omics and non-omics factors conditional on case-control status. The new method provides accurate control of Type I error rate and has increased efficiency over prospective approaches in both simulated and real data.